# **Amendments to the Claims:**

This listing of claims will replace all prior versions, and listings of claims in the application:

# **Listing of Claims:**

1. (Currently Amended): A composition for modulation of LXR function in a eell, said composition comprising a pharmaceutically acceptable excipient and a compound having the formula:

$$A \stackrel{O}{\underset{R^2}{\bigvee}} R^1$$

or a pharmaceutically acceptable salt thereof, wherein

A is <u>adamantyl</u> a member selected from the group consisting of  $(C_5-C_{18})$  alkyl and  $(C_5-C_{18})$  heteroalkyl;

 $R^1$  is a member selected from the group consisting of <u>heteroaryl 1-(furan 2-yl)ethyl, 1-(pyridin-2-yl)ethyl, 1-(furan-2-yl) 2-propyl, 1-(2-pyridyl) 2-propyl, 1-(furanyl)isobutyl, 1-(3-pyridyl)isobutyl, 1-(pyridin-4-yl)ethyl, 1-(pyridin-4-yl)isobutyl, and 1-(3-furanyl) 3-butenyl-heteroaryl( $C_1$ - $C_8$ )alkyl; and</u>

 $R^2$  is a member selected from the group consisting of aryl, heteroaryl when  $R^1$  is heteroaryl and is, aryl( $C_1$ - $C_8$ )alkyl, heteroaryl( $C_1$ - $C_8$ )alkyl when  $R^1$  is heteroaryl( $C_1$ - $C_8$ )alkyl aryl( $C_2$ - $C_8$ )heteroalkyl and heteroaryl( $C_2$ - $C_8$ )heteroalkyl; and said compound binds to the ligand binding domain of LXR $\alpha$  with an affinity of at least 1 micromolar.

- 2. (Currently Amended) A composition in accordance with claim 1, wherein  $\underline{R}^1$  A is selected from the group consisting of 1-(furan-2-yl)ethyl, 1-(pyridin-2-yl)ethyl, 1-(furan-2-yl)-2-propyl, 1-(2-pyridyl)-2-propyl, 1-(furanyl)isobutyl, 1-(3-pyridyl)isobutyl, 1-(pyridin-4-yl)ethyl, 1-(pyridin-4-yl)isobutyl, 1-(2-furanyl)-3-butenyl, and 1-(3-furanyl)-3-butenyl-( $C_5$ - $C_{18}$ )eyeloalkyl and ( $C_5$ - $C_{18}$ )heterocycloalkyl.
  - 3. (Canceled)
  - 4. (Canceled)

- 5. (Currently Amended): A composition in accordance with claim 2[[3]], wherein  $R^1$  is selected from the group consisting of 1-(furan-2-yl)ethyl 1-(3-furanyl)-3-butenyl and 1-(pyridin-2-yl)ethyl.
- 6. (Currently Amended): A composition in accordance with claim  $\underline{1}[[3]]$ , wherein  $R^2$  is selected from the group consisting of aryl and heteroaryl.
- 7. (Currently Amended): A composition comprising a pharmaceutically acceptable excipient and a compound having the formula:

$$A \stackrel{O}{\underset{R^2}{\bigvee}} R^1$$

or a pharmaceutically acceptable salt thereof, in accordance with claim 1, wherein A is adamantyl,  $R^1$  is selected from the group consisting of 1-(furan-2-yl)ethyl, 1-(pyridin-2-yl)ethyl, 1-(furan-2-yl)-2-propyl, 1-(2-pyridyl)-2-propyl, 1-(furanyl)isobutyl, 1-(3-pyridyl)isobutyl, 1-(pyridin-4-yl)ethyl, [[and]] 1-(pyridin-4-yl)isobutyl, 1-(2-furanyl)-3-butenyl, and 1-(3-furanyl)-3-butenyl; and  $R^2$  is selected from the group consisting of heteroaryl  $C_8$  and heteroaryl.

- 8. (Currently Amended): A composition in accordance with claim 7[[1]], wherein A is adamantyl, R<sup>1</sup> is 1-(2-furanyl)-3-butenyl or and 1-(3-furanyl)-3-butenyl and R<sup>2</sup> is selected from phenyl and pyridyl.
  - 9. (Canceled)
- 10. (Currently Amended): A composition in accordance with claim 7[[1]], wherein A is adamantyl, R<sup>1</sup> is selected from the group consisting of 1-(furan-2-yl)ethyl, 1-(3-furanyl)-3-butenyl and 1-(pyridin-2-yl)ethyl and R<sup>2</sup> is selected from the group consisting of aryl and heteroaryl.
  - 11. (Canceled)
- 12. (Currently Amended): A composition in accordance with claim 1, wherein A is 1 adamantyl and R<sup>2</sup> is selected from the group consisting of pyridyl, phenyl, pyrazinyl, pyrimidinyl, pyridazinyl, thiazolyl and furanyl.
  - 13. (Currently Amended): A compound having the formula:

$$A^{1} \bigwedge_{\substack{N \\ R^{21}}}^{N} R^{11}$$

or a pharmaceutically acceptable salt thereof, wherein

 $A^1$  is <u>adamantyl</u> a member selected from the group consisting of ( $C_5$ - $C_{12}$ )monocycloalkyl, ( $C_5$ - $C_{12}$ )heteromonocycloalkyl, ( $C_8$ - $C_{18}$ )bicycloalkyl, ( $C_8$ - $C_{18}$ )heterobicycloalkyl and ( $C_8$ - $C_{18}$ )heterotricycloalkyl;

 $R^{11}$  is a member selected from the group consisting of <u>heteroaryl</u> 1-(furan-2-yl)ethyl, 1-(pyridin-2-yl)ethyl, 1-(furan-2-yl) 2-propyl, 1-(2-pyridyl) 2-propyl, 1-(furanyl)isobutyl, 1-(3-pyridyl)isobutyl, 1-(pyridin-4-yl)ethyl, 1-(pyridin-4-yl)isobutyl, and 1-(3-furanyl) 3-butenyl heteroaryl( $C_1$ - $C_8$ )alkyl; and

 $R^{21}$  is a member selected from the group consisting of aryl, heteroaryl when  $R^1$  is heteroaryl and is, aryl( $C_1$ - $C_8$ )alkyl, heteroaryl( $C_1$ - $C_8$ )alkyl when  $R^1$  is heteroaryl( $C_1$ - $C_8$ )alkyl aryl( $C_2$ - $C_8$ )heteroalkyl and heteroaryl( $C_2$ - $C_8$ )heteroalkyl.

14. (Currently Amended): A compound in accordance with claim 13, wherein  $R^{11}[[A^1]]$  is selected from the group consisting of 1-(furan-2-yl)ethyl, 1-(pyridin-2-yl)ethyl, 1-(pyridin-2-yl)ethyl, 1-(1-(pyridin-2-yl)ethyl, 1-(1-(py

# 15. (Canceled)

- 16. (Previously presented): A compound of claim 13, wherein R<sup>11</sup> is selected from the group consisting of 1-(furan-2-yl)ethyl, 1-(pyridin-2-yl)ethyl and 1-(3-furanyl)-3-butenyl.
- 17. (Currently Amended): A compound in accordance with claim 13, wherein R<sup>21</sup> is selected from the group consisting of aryl and heteroaryl.
  - 18. (Currently Amended): A compound having the formula:

or a pharmaceutically acceptable salt thereof, in accordance with claim 13, wherein  $A^1$  is adamantyl,  $R^{11}$  is selected from the group consisting of 1-(furan-2-yl)ethyl, 1-(pyridin-2-yl)ethyl and 1-(3-furanyl)-3-butenyl, 1-(furan-2-yl)-2-propyl, 1-(2-pyridyl)-2-propyl, 1-(furanyl)isobutyl, 1-(3-pyridyl)isobutyl, 1-(pyridin-4-yl)ethyl, 1-(pyridin-4-yl)isobutyl, 1-(2-furanyl)-3-butenyl, and 1-(3-furanyl)-3-butenyl and  $R^{21}$  is selected from the group consisting of heteroaryl( $C_1$ - $C_8$ )alkyl and heteroaryl.

19. (Currently Amended): A compound in accordance with claim 18[13], wherein  $A^{1}$  is adamantyl,  $R^{11}$  is selected from the group consisting of 1-(2-furanyl)-3-butenyl and 1-(3-furanyl)-3-butenyl and  $R^{21}$  is selected from the group consisting of phenyl and pyridyl.

#### 20. (Canceled)

21. (Currently Amended): A compound in accordance with claim 18[[13]], wherein  $A^{\frac{1}{1}}$  is selected from the group consisting of 1-(furan-2-yl)ethyl, 1-(pyridin-2-yl)ethyl and 1-(3-furanyl)-3-butenyl and  $R^{21}$  is selected from the group consisting of aryl and heteroaryl.

## 22. (Canceled)

23. (Currently Amended): A compound in accordance with claim 18[[13]], wherein  $A^{1}$  is 1-adamantyl,  $R^{11}$  is selected from the group consisting of 1-(furan-2-yl)ethyl, 1-(pyridin-2-yl)ethyl and 1-(3-furanyl)-3-butenyl and  $R^{21}$  is selected from the group consisting of pyridyl, phenyl, pyrazinyl, pyrimidinyl, pyridazinyl, thiazolyl and furanyl.

### 24-27. (Canceled)

- 28. (New): A composition in accordance with claim 7, wherein R<sup>1</sup> is selected from the group consisting of 1-(furan-2-yl)ethyl, 1-(3-furanyl)-3-butenyl and 1-(pyridin-2-yl)ethyl.
  - 29. (New): A composition in accordance with claim 7, wherein R<sup>2</sup> is heteroaryl.
  - 30. (New): A composition in accordance with claim 1, wherein R<sup>2</sup> is pyridyl.
- 31. (New): A composition in accordance with claim 7, wherein R<sup>2</sup> is selected from the group consisting of pyridyl, pyrazinyl, pyrimidinyl, pyridazinyl, thienyl, thiazolyl and furanyl.

- 32. (New): A compound in accordance with claim 18, wherein R<sup>11</sup> is selected from the group consisting of 1-(furan-2-yl)ethyl and 1-(pyridin-2-yl)ethyl and 1-(3-furanyl)-3-butenyl
  - 33. (New): A compound in accordance with claim 18, wherein R<sup>12</sup> is heteroaryl.
- 34. (New): A compound in accordance with claim 13, wherein R<sup>12</sup> is selected from the group consisting of pyridyl, pyrazinyl, pyrimidinyl, pyridazinyl, thienyl, thiazolyl and furanyl.
  - 35. (New): A compound in accordance with claim 13, wherein R<sup>12</sup> is pyridyl.